PIRE LABS BS Certificate of Analysis Registration Certificate Identification Number: 00000007LCND00859758

Next Level Products, LLC

8616 W. Paradise Ln. Peoria, AZ 85382 scott1@gonextlevelup.com (623) 696-1243 Lic. #:

Sample: 2208PURE0182.1157

Strain: Ninja Goldfish Plant Wash RTU

Batch#: NIN-GOLD-1002-01; Batch Size: g; Initial Weight with Packaging: 159.4g Sample Received: 08/08/2022; Report Created: 08/15/2022; Expires: 08/15/2023 Sampled By:

Sample Collection Date and Time:

Ninja Goldfish Plant Wash RTU

Other. Other Lot ID:

Microbials

Pass





BACTERIA (STATE-MANDATED)	Result	Status	Qualifier	BACTERIA (ELECTIVE)	Result Qualifier
E. Coli	Not Detected in	Pass		Listeria	NR
	100 CFU/g			Pseudomonas aeruginosa	NR
Salmonella	Not Detected in 1g	Pass		Staphylococcus aureus	NR
				Patogenic E. coli - STX 1 gene	Not Detected in 1g
				Pathogenic E. coli - STX 2 gene	Not Detected in 1g
FUNGI (STATE-MANDATED)	Result	Status	Qualifier		
Aspergillus flavus	Not Detected in 1g	Pass			
Aspergillus fumigatus	Not Detected in 1g	Pass			
Aspergillus niger	Not Detected in 1g	Pass			
Aspergillus terreus	Not Detected in 1g	Pass			

Date Tested: 08/11/2022 12:00 amNR = No Result; NT = Not Tested; Unless otherwise stated all quality control samples performed within specifications established by the Laboratory

4165 W Clarendon Avenue Phoenix, AZ (623) 334-9194 https://purelabsaz.com/ Lic# 0000007LCND00859758





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This product has been tested by Pure Labs using valid testing methodologies and a quality system as required by state law. Values reported relate only to the product tested. Pure Labs makes no claims as to the efficacy, safety or other risks associated with any detected or non-detected levels of any compounds reported herein. This Certificate shall not be reproduced except in full, without the written approval of Pure Labs. Results apply to the sample as received. Q3 Testing results is for informational purposes only and cannot be used to satisfy dispensary testing requirements in R 9-17-317.01(A) or labeling requirements in R9-17-317.



CSO/QAM

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Qualifiers	Definition
B1	For potency testing, is below the limit of quantitation
B2	When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, is below the maximum allowable concentration in Table 3.1 for the analyte
D1	The limit of quantitation and the sample results were adjusted to reflect sample dilution
11	The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance criteria in subsection (L)(1) with respect to the reference spectra, indicating interference –
L1	When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, the percent recovery of a laboratory control sample is greater than the acceptance limits in subsection (K)(2)(c), but the sample's target analytes were not detected above the maximum allowable concentrations in Table 3.1 for the analytes in the sample
M1	High, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
M2	Low, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
М3	Unusable because the analyte concentration was disproportionate to the spike level, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
M4	The analysis of a spiked sample required a dilution such that the spike recovery calculation does not provide useful information, but the recovery from the associated laboratory control sample in subsection $(K)(2)$ was within acceptance criteria
M5	The analyte concentration was determined by the method of standard addition, in which the standard is added directly to the aliquots of the analyzed sample
N1	A description of the variance is described in the final report of testing according to R9-17- 404.06(B)(3)(d)(ii)
R1	404.06(B)(3)(d)(ii) – N1; 9. The relative percent difference for the laboratory control sample and duplicate exceeded the limit in subsection (K)(3), but the recovery in subsection (K)(2) was within acceptance criteria
R2	The relative percent difference for a sample and duplicate exceeded the limit in subsection (O)
V1	The recovery from continuing calibration verification standards exceeded the acceptance limits in subsection $(J)(1)(b)$, but the sample's target analytes were not detected above the maximum allowable concentrations in Table 3.1 for the analytes in the sample
Q1	Sample integrity was not maintained
Q2	The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices
Q3	Testing result is for informational purposes only and cannot be used to satisfy dispensary testing requirements in R9-17-317.01(A) or labeling requirements in R9-17-317
ND	Analyte not detected @ / or above the reporting limit
RPD	Relative % difference
%REC	% recovery
Source	Sample that was matrix spiked or duplicated.

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RIRE LABS



Craig Knoblock CSO/QAM

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